

Eylor
08/644289

08/644289

FILE 'REGISTRY' ENTERED AT 15:48:13 ON 24 JUN 1999

L12 1 S SLRPFKALVREKGRPSHSC/SQSP

=> d .bevreg1

L12 ANSWER 1 OF 1 REGISTRY COPYRIGHT 1999 ACS

RN 173787-20-7 REGISTRY

CN L-Cysteine, L-seryl-L-leucyl-L-arginyl-L-prolyl-L-phenylalanyl-L-lysyl-L-alanyl-L-leucyl-L-valyl-L-arginyl-L-.alpha.-glutamyl-L-lysylglycyl-L-histidyl-L-arginyl-L-prolyl-L-seryl-L-histidyl-L-seryl-
(9CI) (CA INDEX NAME)

SQL 20

SEQ 1 SLRPFKALVR EKGHRPSHSC

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HITS AT: 1-20

REFERENCE 1: 128:216370

REFERENCE 2: 128:21873

REFERENCE 3: 124:173450

(FILE 'CAPLUS' ENTERED AT 15:49:12 ON 24 JUN 1999)

L13 3 S L12

=> d 1-3 .bevstr

L13 ANSWER 1 OF 3 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1998:178120 CAPLUS

DOCUMENT NUMBER: 128:216370

TITLE: Recombinant p53as protein and antibody for diagnosis and therapy of malignancy

INVENTOR(S): Kulesz-Martin, Molly F.

PATENT ASSIGNEE(S): Health Research, Inc., USA

SOURCE: U.S., 30 pp. Cont.-in-part of U.S. Ser. No. 259,612.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 6

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5726024	A	19980310	US 96-644291	19960510
US 5688918	A	19971118	US 94-259612	19940614
EP 806478	A2	19971112	EP 97-107696	19970512
EP 806478	A3	19990609		

Searcher : Shears 308-4994

R: BE, CH, DE, DK, FR, GB, LI, NL, SE

PRIORITY APPLN. INFO.: US 93-100496 19930802
 US 94-195952 19940211
 US 94-259612 19940614
 US 96-644289 19960510
 US 96-644291 19960510
 US 96-644456 19960510

AB The invention comprises plasmids and viral vectors contg. an animal p53as (alternatively spliced p53) cDNA sequence. A portion of the p53as sequence may be identified to a position of wild type p53 gene from the same animal. In preferred embodiments, the p53as is mouse or human p53as. A preferred viral vector is baculovirus vector. The invention further includes antibodies both polyclonal and monoclonal, to p53as and to at least a portion of human p53 intron 10 sequence encoding SLRPFKALVREKGRPSSHSC, which is related to p53as sequences and plasmids and viral vectors contg. such sequences. All of the above find utility in studying p53 and p53as and their relative expressions which is believed important for detection and control of malignant cells and their susceptibility to treatment agents. The antibodies can detect the presence of p53as and related sequences and when injected into cells could cause cell cycle arrest and the plasmids and viral vectors, with appropriate promoters, can cause expression of the p53as and p53 intron 10 sequences which can affect cell growth and perhaps arrest certain malignancies.

IT 173787-20-7

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (recombinant p53as protein and antibody for diagnosis and therapy of malignancy)

L13 ANSWER 2 OF 3 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1997:761610 CAPLUS
 DOCUMENT NUMBER: 128:21873
 TITLE: P53as protein and antibody therefor
 INVENTOR(S): Kulesz-Martin, Molly F.
 PATENT ASSIGNEE(S): Health Research, Inc., USA
 SOURCE: U.S., 25 pp. Cont.-in-part of U.S. Ser. No. 195,952.
 CODEN: USXXAM
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 6
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5688918	A	19971118	US 94-259612	19940614
EP 652232	A1	19950510	EP 94-610042	19940801

Searcher : Shears 308-4994

08/644289

R: BE, CH, DE, DK, FR, GB, LI, NL, SE

JP 08099998	A2	19960416	JP 94-181558	19940802
CA 2150994	AA	19951215	CA 95-2150994	19950605
JP 08081500	A2	19960326	JP 95-169323	19950613
EP 709397	A1	19960501	EP 95-610034	19950614

R: BE, CH, DE, DK, FR, GB, LI, NL, SE

US 5726024	A	19980310	US 96-644291	19960510
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PRIORITY APPLN. INFO.:

US 93-100496	19930802
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US 94-195952	19940211
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US 94-259612	19940614
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AB The invention comprises plasmids and viral vectors contg. an animal p53as cDNA sequence. A portion of the p53as sequence may be identified to a position of wild type p53 gene from the same animal. In preferred embodiments, the p53as is mouse or human p53as. A preferred viral vector is baculovirus vector. The invention further includes antibodies both polyclonal and monoclonal, to p53as and to at least a portion of human p53 intron 10 sequence encoding SLRPFKALVREKGRPSSHSC which is related to p53as sequences and plasmids and viral vectors contg. such sequences. All of the above find utility in studying p53 and p53as and their relative expressions which is believed important for detection and control of malignant cells and their susceptibility to treatment agents. The antibodies can detect the presence of p53as and related sequences and when injected into cells could cause cell cycle arrest and the plasmids and viral vectors, with appropriate promoters, can cause expression of the p53as and p53 intron 10 sequences which can affect cell growth and perhaps arrest certain malignancies.

IT 173787-20-7

RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(recombinant p53as protein and antibody for diagnosis and therapy of malignancy)

L13 ANSWER 3 OF 3 CAPLUS COPYRIGHT 1999 ACS

ACCESSION NUMBER: 1996:126792 CAPLUS

DOCUMENT NUMBER: 124:173450

TITLE: Monoclonal and polyclonal antibody to alternatively spliced p53 protein (p53as) for diagnosis and prognosis of cancer

INVENTOR(S): Kulesz-Martin, Molly F.

PATENT ASSIGNEE(S): Health Research, Inc., USA

SOURCE: Can. Pat. Appl., 40 pp.

CODEN: CPXXEB

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 6

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	Searcher	:	Shears	308-4994

08/644289

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CA 2150994	AA	19951215	CA 95-2150994	19950605
US 5688918	A	19971118	US 94-259612	19940614
PRIORITY APPLN. INFO.:			US 94-259612	19940614
			US 93-100496	19930802
			US 94-195952	19940211

AB The invention comprises plasmids and viral vectors contg. an animal p53as cDNA sequence. A portion of the p53 sequence may be identified to a position of wild type p53 gene from the same animal. In preferred embodiments, the p53as is mouse or human p53as. A preferred viral vector is baculovirus vector. The invention further includes antibodies both polyclonal and monoclonal, to p53as and to at least a portion of human p53 intron 10 sequence encoding SLRPFKALVREKGHRPSHSC which is related to p53as sequences and plasmids and viral vectors contg. such sequences. All of the above find utility in studying p53 and p53as and their relative expressions which is believed important for detection and control of malignant cells and their susceptibility to treatment agents. The antibodies can detect the presence of p53as and related sequences and when injected into cells could cause cell cycle arrest and the plasmids and viral vectors, with appropriate promoters, can cause expression of the p53as and p53 intron 10 sequences which can affect cell growth and perhaps arrest certain malignancies.

IT 173787-20-7

RL: BSU (Biological study, unclassified); BIOL (Biological study)
(polyclonal or monoclonal antibody to human or mouse p53as
proteins for diagnosis or prognosis of cancer)

=> fil hom

FILE 'HOME' ENTERED AT 15:49:47 ON 24 JUN 1999

Searcher : Shears 308-4994

 MWSEKELI
 (TM)

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 Distribution rights by Oxford Molecular Ltd

Mparch.p protein - protein database search, using Smith-Waterman algorithm

Run on: Wed Jun 23 14:53:55 1999; Maspar time 41.91 Seconds

Tabular output not generated. 10.148 Million cell updates/sec

Title: >US-08-644-289-1

Description: (1-20) from US08644289 .pep

Sequence: 1 SLRPFKALVREKGRHPSHC 20

Scoring table:

PAM 150
 Gap 15

Searched: 170751 seqs, 2126608 residues

Post-processing: Minimum Match 0%

Listing first 1000 summaries
 Maximum DB seq length 30

Database:

a:geneseqs
 1:part1.2:part2.3:part3.4:part4.5:part5.6:part6.7:part7
 8:part8.9:part9.10:part10.11:part11.12:part12.13:part13
 14:part14.15:part15.16:part16.17:part17.18:part18
 19:part19.20:part20.21:part21.22:part22.23:part23
 24:part24.25:part25.26:part26.27:part27.28:part28
 29:part29.30:part30.31:part31.32:part32.33:part33
 34:part34.35:part35.36:part36.37:part37.38:part38
 39:part39

Statistics: Mean 20.743; Variance 62.468; scale 0.332

Pred. No. is the number of results predicted by chance to have a
 score greater than or equal to the score of the result being printed,
 and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description	Pred. No.
1	151	100.0	20	18	R92698 Human wild-type p53 g	5.67e-10
2	136	90.1	21	29	W33235 Human p53 fragment.	4.52e-08
3	85	56.3	20	29	W33239 Peptide of the human	7.01e-02
4	50	33.1	18	8	R44726 HEV ORF1 peptide (123	4.09e+02
5	48	31.8	17	29	W33236 Mouse p53as peptide.	6.37e+02
6	47	31.1	24	7	R37861 HIV protein fragment	7.92e+02
7	46	30.5	16	9	R45673 RNP heparin binding	9.83e+02
8	46	30.5	20	26	W26230 pMA2-T8-19g junction	9.83e+02
9	45	29.8	29	9	R45671 N terminus of human C	9.83e+02
10	45	29.8	14	6	R32978 Mastoparan analogue (1.22e+03
11	45	29.8	19	24	W28500 Last 19 C-terminal am	1.22e+03
12	45	29.8	27	3	R11996 N-terminal of p29 pro	1.22e+03
13	44	29.1	15	21	W15386 N-alpha-Ac-(Ala26,Phe	1.50e+03
14	44	29.1	15	20	W12816 Peptide chain ENCFE6V	1.50e+03
15	44	29.1	15	33	W51822 Peptide YX analogue #	1.50e+03

Note: Post-processor removed 968 summaries from list due to search parameters chosen.

ALIGNMENTS

RESULT ID	Score	Query Match	Length	ID	Description	Pred. No.
1	151	100.0	20	18	R92698 Human wild-type p53 gene Intron 10 encoded epitope.	5.67e-10
2	136	90.1	21	29	W33235 Human p53 fragment.	4.52e-08
3	85	56.3	20	29	W33239 Peptide of the human	7.01e-02
4	50	33.1	18	8	R44726 HEV ORF1 peptide (123	4.09e+02
5	48	31.8	17	29	W33236 Mouse p53as peptide.	6.37e+02
6	47	31.1	24	7	R37861 HIV protein fragment	7.92e+02
7	46	30.5	16	9	R45673 RNP heparin binding	9.83e+02
8	46	30.5	20	26	W26230 pMA2-T8-19g junction	9.83e+02
9	45	29.8	29	9	R45671 N terminus of human C	9.83e+02
10	45	29.8	14	6	R32978 Mastoparan analogue (1.22e+03
11	45	29.8	19	24	W28500 Last 19 C-terminal am	1.22e+03
12	45	29.8	27	3	R11996 N-terminal of p29 pro	1.22e+03
13	44	29.1	15	21	W15386 N-alpha-Ac-(Ala26,Phe	1.50e+03
14	44	29.1	15	20	W12816 Peptide chain ENCFE6V	1.50e+03
15	44	29.1	15	33	W51822 Peptide YX analogue #	1.50e+03

PR 10-MAY-1996; US-644289.
PR 10-MAY-1996; US-644291.
PA (HEAL-) HEALTH RES INC.
PI Kulesz-Martin MF;
DR WPI: 97-538617/50.
PT Alternatively spliced synthetic p53 protein, p53as - useful to
PT generate specific antibody for cell growth and differentiation
PT research
PS Disclosure: Page 3; 67pp; English.
CC This peptide is a fragment of the human p53 protein. It has at least
CC 80% homology with a novel p53as protein for mammals, and the same
CC sequence specific binding on the cellular level as active p53 protein
CC from that mammal, except that the sequence specific binding of the novel
CC peptide remains active in cellular environments in which the sequence
CC specific binding of p53 protein is deactivated. The novel peptide
CC differs from p53 protein in the final 50 carboxy-terminal amino acids of
CC p53 protein, and contains a specific antigen site not present in p53
CC protein, giving rise to an antibody unique for the p53as peptide.
CC Antibodies specific for murine p53as protein can be used in cell growth
CC and differentiation basic research. The development of a homologous
CC protein for use in human cells, would have applications in the diagnosis
CC and prognosis of human diseases such as cancer, and in the design of
CC treatment strategies for such diseases. The association of p53as protein
CC expression with the G2 phase of the cell cycle suggests a functional role
CC in G2 arrest, and the potential use of the p53as coding sequence for gene
CC therapy.
SQ Sequence 21 AA:

Query Match 90.1%; Score 136; DB 29; Length 21;
Best Local Similarity 95.2%; Pred. No. 4.52e-08;
Matches 20; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Db 1 slrpfkalvrekhpshsc 21
|||
Qy 1 SLRPFKALVREKGRPS-HSC 20

RESULT 3
ID W33239 standard; peptide; 20 AA.
AC W33239;
DT 28-MAY-1998 (first entry)
DE Peptide of the human p53as protein.
KW Human p53as peptide; mouse p53 protein; antigen; antibody; cell growth;
KM differentiation; cancer; G2 phase; gene therapy.
OS Homo sapiens.
PN EP-806478-A2.
PD 12-NOV-1997.
PF 12-MAY-1997; 107696.
PR 10-MAY-1996; US-644456.
PR 10-MAY-1996; US-644289.
PR 10-MAY-1996; US-644291.
PA (HEAL-) HEALTH RES INC.
PI Kulesz-Martin MF;
DR WPI: 97-538617/50.
PT Alternatively spliced synthetic p53 protein, p53as - useful to
PT generate specific antibody for cell growth and differentiation
PT research
PS Disclosure: Page 11; 67pp; English.
CC This peptide is unique to human p53as, and is encoded by intron 10. It
CC was used to immunize rabbit in order to generate polyclonal antibodies
CC to mouse p53as. The novel peptide (p53as) differs from p53 protein
CC in the final 50 carboxy-terminal amino acids of p53 protein, and
CC contains a specific antigen site not present in p53 protein, giving rise
CC to an antibody unique for the p53as peptide. Antibodies specific for
CC murine p53as protein can be used in cell growth and differentiation basic
CC research. The development of a homologous protein for use in human
CC cells, would have applications in the diagnosis and prognosis of human
CC diseases such as cancer, and in the design of treatment strategies for
CC such diseases. The association of p53as protein expression with the G2
CC phase of the cell cycle suggests a functional role in G2 arrest, and the
CC potential use of the p53as coding sequence for gene therapy.
SQ Sequence 20 AA;

Query Match 56.3%; Score 85; DB 29; Length 20;
Best Local Similarity 100.0%; Pred. No. 7.01e-02;
Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 rekghrpshsc 11
|||
Qy 10 REKGRPSHSC 20

RESULT 4
ID R44726 standard; peptide; 18 AA.
AC R44726;
DT 11-JAN-1994 (first entry)
DE HEV ORF1 peptide (1237-1254).
KW Enterically transmitted non-A non-B hepatitis; ET-NANB;
KW vaccine.
OS Hepatitis E virus.
PN WO9314116-A.
PD 22-JUL-1993.
PF 15-JAN-1993; U00459.
PR 17-JAN-1992; US-822335.
PR 01-MAY-1992; US-876941.
PA (GENE-) GENELABS TECHNOLOGIES INC.
PA (USSH) US SEC DEPT HEALTH.
PI Bradley DW, Carl M, Reyes GR, Tam AW;
DR WPI: 93-243144/30.
PT New immunogenic hepatitis E virus (HEV) peptide(s) - are from the
PT ORF1, ORF2 and ORF3 regions of HEV, useful as a vaccine against
PT HEV infection.
PS Claim 1, Fig 11; 48pp; English.
CC Immunogenic hepatitis E virus (HEV) peptides are selected from the
CC ORF1, ORF2 and ORF3 regions of HEV. The peptides can be used in
CC vaccines to prevent infection by HEV. The antibodies can neutralise
CC and block HEV infection and can be used to prevent or treat HEV
CC infection. The peptides and antibodies can also be used as
CC diagnostic reagents.
SQ Sequence 18 AA;

Query Match 33.1%; Score 50; DB 8; Length 18;
Best Local Similarity 46.7%; Pred. No. 4.09e-02;
Matches 7; Conservative 2; Mismatches 6; Indels 0; Gaps 0;

Db 3 lsafhglaelghrp 17
|||
Qy 2 LRPFKALVREKGRHP 16

RESULT 5
ID W33236 standard; peptide; 17 AA.
AC W33236;
DT 28-MAY-1998 (first entry)
DE Mouse p53as peptide.
KW Mouse p53as peptide; human p53 protein; antigen; antibody; cell growth;
KM differentiation; cancer; G2 phase; gene therapy.
OS Mus sp.
PN EP-806478-A2.
PD 12-NOV-1997.
PF 12-MAY-1997; 107696.
PR 10-MAY-1996; US-644456.
PR 10-MAY-1996; US-644289.
PR 10-MAY-1996; US-644291.
PA (HEAL-) HEALTH RES INC.
PI Kulesz-Martin MF;
DR WPI: 97-538617/50.
PT Alternatively spliced synthetic p53 protein, p53as - useful to
PT generate specific antibody for cell growth and differentiation
PT research
PS Disclosure: Page 7; 67pp; English.
CC This is a peptide from an alternatively spliced mouse p53as gene. It
CC has at least 80% homology with a fragment of the human p53 protein, and
CC the same sequence specific binding on the cellular level as active p53
CC protein from that mammal, except that the sequence specific binding of
CC the novel peptide remains active in cellular environments in which the

CC sequence specific binding of p53 protein is deactivated. The novel
 CC peptide differs from p53 protein in the final 50 carboxy-terminal amino
 CC acids of p53 protein, and contains a specific antigen site not present
 CC in p53 protein, giving rise to an antibody unique for the p53as peptide.
 CC Antibodies specific for murine p53as protein can be used in cell growth
 CC and differentiation basic research. The development of a homologous
 CC protein for use in human cells, would have applications in the diagnosis
 CC and prognosis of human diseases such as cancer, and in the design of
 CC treatment strategies for such diseases. The association of p53as protein
 CC expression with the G2 phase of the cell cycle suggests a functional role
 CC in G2 arrest, and the potential use of the p53as coding sequence for gene
 CC therapy.
 SQ Sequence 17 AA;

Query Match 31.8%; Score 48; DB 29; Length 17;
 Best Local Similarity 55.6%; Pred. No. 6.37e+02;
 Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Db 4 rafqalike 12
 :|:|:|:|:
 Oy 3 RPFKALVRE 11

RESULT 6
 ID R37861 standard; peptide; 24 AA.
 AC R37861;
 DT 21-OCT-1993 (first entry)
 DE HIV protein fragment vif-C(170-192).
 KW Human immunodeficiency virus; screening; detection; AIDS;
 KM acquired immune deficiency syndrome; viral infectivity.
 OS Synthetic.
 FH Key Location/Qualifiers
 FT misc_difference 1 /note="opt.absent"
 FN J05125096-A.
 PD 21-MAY-1993;
 PR 02-NOV-1991; 288159.
 PA (YAMH) NIPPON STEEL CHEM CO.
 PA (YAMA) NIPPON STEEL CORP.
 DR WPI: 93-200512/25.
 PT New peptide(s) - useful for diagnosis of HIV infection
 PS Claim 1; Page 2; 11pp; Japanese.
 CC This peptide has a sequence corresponding to part of the HIV
 CC C-terminal region. It is useful as a diagnostic agent for screening
 CC and monitoring HIV carriers. See also R37860.
 SQ Sequence 24 AA;

Query Match 31.1%; Score 47; DB 7; Length 24;
 Best Local Similarity 60.0%; Pred. No. 7.92e+02;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Db 11 kckghrsht 20
 :|:|:|:|:
 Oy 10 REKGRHRSMS 19

RESULT 7
 ID R45673 standard; Protein; 16 AA.
 AC R45673;
 DT 25-JUL-1994 (first entry)
 DE RNIP heparin binding fragment.
 KW Cationic antibacterial protein; lipopolysaccharide binding;
 KM anticoagulant; granulocytes; RNIP; LPS; sepsis; autoimmune disorder;
 OS septic shock; rabbit; CAP18.
 OS Synthetic.
 PN W09402589-A.
 PD 03-FEB-1994.
 PF 15-JUL-1993; U06731.
 PR 17-JUL-1992; US-916761.
 PR 17-JUL-1992; US-916765.
 PA (PANO-) PANORAMA RES INC.
 PI Hirata M, Larrick JW, Wright SC;

DR WPI: 94-048847/06.
 PT Sequences encoding mammalian cationic antibacterial proteins -
 PT are homologous to human and rabbit CAP18 sequences and have
 PT lipopolysaccharide binding and anti-coagulation activity
 PS Disclosure; Page 50; 112pp; English.
 CC The sequence of CAP18 C-terminal RNIP was compared to that of a
 CC number of heparin binding proteins to determine residues important for
 CC binding to lipopolysaccharides and inhibiting LPS-mediated activation
 CC of macrophage, as well as interfering with the clotting cascade to
 CC inhibit coagulation in conditions of disseminated intravascular
 CC coagulation.
 CC See also R45667-81.
 SQ Sequence 16 AA;

Query Match 30.5%; Score 46; DB 9; Length 16;
 Best Local Similarity 45.5%; Pred. No. 9.83e+02;
 Matches 5; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Db 6 lkfrnklike 16
 :|:|:|:|:
 Oy 2 LRFKALVREK 12

RESULT 8
 ID W26230 standard; peptide; 20 AA.
 AC W26230;
 DT 16-MAR-1998 (first entry)
 DE pMA2-TH-IgG junction region of fusion proteins.
 KW Fusion protein; hydrophilic spacer; recombinant; expression system;
 KM carboxypeptidase.
 OS Synthetic.
 PN W09728272-A1.
 PD 07-AUG-1997.
 PR 31-JAN-1997;
 PR 31-JAN-1996; US-595043.
 PA (TECH-) TECHNOLOGENE INC.
 PI Sgarlato GD;
 DR WPI: 97-402624/37.
 DR N-PSDB: R80156.
 PT Recombinant protein expression system for fusion protein production
 PT - useful for high quantity production of authentic recombinant
 PT proteins
 PS Example 1; Fig 7; 194pp; English.
 CC A novel recombinant vector has been developed which comprises a
 CC nucleotide sequence encoding a fusion protein. The fusion protein
 CC comprises three domains joined together in order, from N-terminus to
 CC C-terminus, of a first domain comprising a protein of interest, a second
 CC domain comprising a hydrophilic spacer and an affinity domain, each
 CC domain comprising amino acid residues. The present sequence represents
 CC a junction region (i.e. the region which joins the protein of interest
 CC with the affinity domain) present in pMA2-TH-IgG, used in example 1
 CC of the present invention. The recombinant vector is used for the
 CC production of authentic recombinant proteins of interest. The method of
 CC the invention is useful for the expression of fusion proteins capable of
 CC isolation by affinity chromatography in pro- or eukaryotic cells. This
 CC method allows for the efficient cleavage and generation of authentic
 CC proteins of interest that do not contain extraneous (i.e. non-naturally
 CC occurring) amino acids.
 SQ Sequence 20 AA;

Query Match 30.5%; Score 46; DB 26; Length 20;
 Best Local Similarity 35.3%; Pred. No. 9.83e+02;
 Matches 6; Conservative 5; Mismatches 5; Indels 1; Gaps 1;

Db 4 sfrrlvp-rgrrtcppc 19
 :|:|:|:|:|:|:
 Oy 4 PFKALVREKGRHRSMS 20

RESULT 9
 ID R45671 standard; Protein; 29 AA.
 AC R45671;
 DT 25-JUL-1994 (first entry)

DE N terminus of human CAP18.
 KM Cationic antibacterial protein; lipopolysaccharide binding;
 KM anticoagulant; granulocytes; RNIP; LPS; sepsis; autoimmune disorder;
 KM septic shock; rabbit.
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT misc-difference 23 /label= Asp, Lys
 FT misc-difference 26 /label= Gln, Ile
 FT misc-difference 27 /label= Gly, Gln
 FT
 PN MO9402589-A.
 PD 03-FEB-1994.
 PE 15-JUL-1993: US-06731.
 PR 17-JUL-1992: US-916761.
 PR 17-JUL-1992: US-916765.
 PA (PANO-) PANORAMA RES INC.
 PI Hirata M, Larrick JW, Wright SC;
 DR WPI: 94-048847/06.
 PT Sequences encoding mammalian cationic antibacterial proteins -
 PT are homologous to human and rabbit CAP18 sequences and have
 PT Lipo:polysaccharide binding and anti-coagulation activity
 PS Disclosure: Page 59; 112pp; English.
 CC The sequence is that of a human cationic antibacterial protein CAP18
 CC N-terminal fragment obtd. from granulocytes as sequenced. The
 CC sequence corresponds to the N-terminus of the C-terminal RNIP fragment
 CC of rabbit CAP18. The fragment is capable of binding to lipopolysaccharide
 CC and inhibiting LPS-mediated activation of macrophage, as well as
 CC interfering with the clotting cascade to inhibit coagulation in
 CC conditions of disseminated intravascular coagulation. The polypeptides
 CC can also be used to attenuate, inhibit or prevent LPS-associated
 CC conditions, e.g. sepsis, autoimmune disorders, inflammation, etc.
 CC See also R45667-81.
 SQ Sequence 29 AA;
 QY
 DB 6 lrfkrnkkek 16
 QY 2 LRPFKALVREK 12
 RESULT 10
 ID R32978 standard; peptide: 14 AA.
 AC R32978;
 DT 02-JUL-1993 (first entry)
 DE Mastoparan analogue (41) having G protein modulatory activity.
 KM Mastoparan; MP; cellular regulation; receptor-based analogue;
 KM G protein; guanine nucleotide binding regulatory protein;
 KM asthma; ulcer; cardiovascular disease; allergy; Parkinson's;
 KM small cell carcinoma; lung; glaucoma; respiratory tract congestion;
 KM inflammation.
 OS Synthetic.
 PN MO9303749-A.
 PD 04-MAR-1993.
 PE 14-AUG-1992: US-06825.
 PR 21-AUG-1991: US-748319.
 PA (TEXA) UNIV TEXAS SYSTEM.
 PI Higashijima T, Ross EM;
 DR WPI: 93-093715/11.
 PT New peptide(s) contg. mastoparan- or receptor-analogue region -
 PT uses as G protein modulators, for treating asthma, ulcers,
 PT cardiovascular disorders and Parkinson's disease
 PS Claim 43; Page 83; 96pp; English.
 CC The peptide is an example of a highly generic formula, and is
 CC represented as found in the disclosure of the specification. The
 CC claimed peptide lacks the N-terminal 19 amino acids.
 CC The peptide is capable of modulating G protein action in a cell. It
 CC may therefore be used for treating diseases involving G proteins,
 CC e.g. asthma, ulcers, cardiovascular diseases, allergies, Parkinson's

CC disease, small cell carcinoma of the lung, glaucoma, respiratory
 CC tract congestion or inflammation.
 SQ Sequence 14 AA;
 QY
 DB 2 nlralfalar 11
 QY 1 LRPFKALVR 10
 RESULT 11
 ID M28500 standard; Protein: 19 AA.
 AC M28500;
 DT 18-NOV-1997 (first entry)
 DE Last 19 C-terminal amino acids of alternatively spliced murine p53.
 KM Leucine zipper domain; LZD; oligomerisation domain; mutant; mutin;
 KM substitution; replacement; anti-oncogene; hyperproliferation;
 KM cancer; restenosis; tumour suppression; apoptosis; AS p53; mouse.
 OS Mus musculus.
 PN MO9704092-A1.
 PD 06-FEB-1997.
 PE 17-JUL-1996: F01111.
 PR 19-JUL-1995: FR-008729.
 PA (RHON) RHONE-POULENC ROBER SA.
 PI Bracco L, Conseiller E;
 DR WPI: 97-13263/12.
 DR N-P5DB; T8621.
 PT New p53 variants e.g. with oligomerisation domain replaced by
 PT leucine zipper - useful for treating hyper-proliferative disorders,
 PT esp. cancer and restenosis
 PS Example A: Page 29; 133pp; French.
 CC A claimed p53 variant consists of a fragment coding for amino
 CC acids 1-366 of human p53 protein, followed by a fragment coding for
 CC the last 19 C-terminal amino acids of the alternatively spliced (AS)
 CC form of murine p53 (i.e. the present sequence). The variant is a
 CC more active and more stable tumour suppressor and apoptosis-inducing
 CC agent than wild-type p53 and is active where the wild-type protein
 CC is not.
 SQ Sequence 19 AA;
 QY
 DB 6 rafqalmke 14
 QY 3 RPFKALVRE 11
 RESULT 12
 ID R1196 standard; peptide: 27 AA.
 AC R1196;
 DT 26-JUL-1991 (first entry)
 DE N-terminal of p29 protein.
 KM Wegener's granulomatosis; monoclonal antibodies; autoantibodies;
 KM glomerulonephritis; serine protease; antigen.
 OS Homo sapiens.
 PN MO9106572-A.
 PD 16-MAY-1991.
 PE 29-OCT-1990: US-06277.
 PR 27-OCT-1989: US-428286.
 PA (GEHO-) GEN HOSPITAL CORP.
 PI Arnaut M, McCluskey RT, Niles J;
 DR WPI: 91-164137/22.
 PT Purified p29 protein - used to detect auto-antibodies diagnostic
 PT for Wegener's granulomatosis and conditions associated with
 PT glomerulo nephritis
 PS Claim 1; Page 22; 33pp; English.
 CC The p29 protein is a 29 kD antigen which was prepd. by affinity
 CC purification. from neutrophil acid extract, using 188 monoclonal

CC are truncated versions of peptide YX. They interact solely with peptide
CC YX receptors and not with homologous receptors such as NPY Y1 and Y3,
CC thus minimizing unwanted (ant)agonist side reactions. The present
CC sequence represents a peptide YX analogue.
SQ Sequence 15 AA:

Query Match 29.1%; Score 44; DB 33; Length 15;
Best Local Similarity 53.8%; Pred. No. 1.50e+03;
Matches 7; Conservative 3; Mismatches 2; Indels 1; Gaps 1;
Db 2 siraflnlvtqr 14
|||:| |||:
QY 1 SIRPFKALV-REK 12

Search completed: Wed Jun 23 14:56:08 1999
Job time : 133 secs.

CC / APPLICATION NUMBER: US/08/644,291
CC FILING DATE: 10-May-1996
CC CLASSIFICATION: 530
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 08/259,612
CC FILING DATE: 14-Jun-1994
CC CLASSIFICATION: 530
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 08/195,952
CC FILING DATE: 11-Feb-1994
CC CLASSIFICATION: 530
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 08/100,496
CC FILING DATE: 02-Aug-1993
CC CLASSIFICATION: 530
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Dunn, Michael L.
CC REGISTRATION NUMBER: 25,330
CC REFERENCE/DOCKET NUMBER: RPP:135E US
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (716) 433-1661
CC TELEFAX: (716) 433-1665
CC INFORMATION FOR SEQ ID NO: 1:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 21
CC TYPE: Amino Acid
CC STRANDEDNESS: Unknown
CC TOPOLOGY: Unknown
CC MOLECULE TYPE: Peptide
CC HYPOTHETICAL: NO
CC ANTI-SENSE: NO
CC FRAGMENT TYPE:
CC ORIGINAL SOURCE:
CC ORGANISM:
CC STRAIN:
CC INDIVIDUAL ISOLATE:
CC DEVELOPMENTAL STAGE:
CC HAPLOTYPE:
CC TISSUE TYPE:
CC CELL TYPE:
CC CELL LINE:
CC ORGANELLE:
CC IMMEDIATE SOURCE:
CC LIBRARY:
CC CLONE:
CC POSITION IN GENOME:
CC CHROMOSOME/SEGMENT: 17
CC MAP POSITION: Intron 10
CC UNITS:
CC FEATURE:
CC NAME/KEY:
CC LOCATION:
CC IDENTIFICATION METHOD:
CC OTHER INFORMATION:
CC PUBLICATION INFORMATION:
CC AUTHORS:
CC TITLE:
CC JOURNAL:
CC VOLUME:
CC ISSUE:
CC PAGES:
CC DATE:
CC DOCUMENT NUMBER:
CC FILING DATE:
CC PUBLICATION DATE:
CC RELEVANT RESIDUES IN SEQ ID NO:
SQ SEQUENCE 21 AA; 2393 MW; 2418 CN;

Query Match 90.1%; Score 136; DB 1; Length 21;
Best Local Similarity 95.2%; Pred. No. 2,71e-08;
Matches 20; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Db 1 SLRPFKALVREKGRPSHSC 21

QY 1 SLRPFKALVREKGRPS-HSC 20
|||||
RESULT 2 STANDARD; PRT; 21 AA.
ID US-08-259-612A-1
AC xxxxxx
XX
XX
DT
DE Sequence 1, Application US/08259612A
XX Sequence 1, Application US/08259612A
CC Patent No. 5688918
CC GENERAL INFORMATION:
CC APPLICANT: Kulacz-Martin, Molly F.
CC TITLE OF INVENTION: p53as PROTEIN AND ANTIBODY
CC TITLE OF INVENTION: THEREFOR
CC NUMBER OF SEQUENCES: 9
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Dunn & Associates, P.C.
CC STREET: P.O. Box 96
CC CITY: Newfane
CC STATE: New York
CC COUNTRY: U.S.A.
CC ZIP: 14108
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Diskette - 3.50 inch, 1.44 MB
CC MEDIUM TYPE: Storage
CC COMPUTER: Victor 300 SX/25 (IBM PC Compatible)
CC OPERATING SYSTEM: MS-DOS Version 5.0
CC SOFTWARE: Wordstar Professional Release 4
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/259,612A
CC FILING DATE: 14-Jun-1994
CC CLASSIFICATION: 530
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 08/195,952
CC FILING DATE: 14-Feb-1994
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Dunn, Michael L.
CC REGISTRATION NUMBER: 25,330
CC REFERENCE/DOCKET NUMBER: RPP:135B US
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (716) 433-1661
CC TELEFAX: (716) 433-1665
CC INFORMATION FOR SEQ ID NO: 1:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 21
CC TYPE: Amino Acid
CC STRANDEDNESS: Unknown
CC TOPOLOGY: Unknown
CC MOLECULE TYPE: Peptide
CC HYPOTHETICAL: NO
CC ANTI-SENSE: NO
CC FRAGMENT TYPE:
CC ORIGINAL SOURCE:
CC ORGANISM:
CC STRAIN:
CC INDIVIDUAL ISOLATE:
CC DEVELOPMENTAL STAGE:
CC HAPLOTYPE:
CC TISSUE TYPE:
CC CELL TYPE:
CC CELL LINE:
CC ORGANELLE:
CC IMMEDIATE SOURCE:
CC LIBRARY:
CC CLONE:
CC POSITION IN GENOME:
CC CHROMOSOME/SEGMENT: 17
CC MAP POSITION: Intron 10

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CC UNITS:
CC FEATURE:
CC NAME/KEY:
CC LOCATION:
CC IDENTIFICATION METHOD:
CC OTHER INFORMATION:
CC PUBLICATION INFORMATION:
CC AUTHORS:
CC TITLE:
CC JOURNAL:
CC VOLUME:
CC ISSUE:
CC PAGES:
CC DATE:
CC DOCUMENT NUMBER:
CC FILING DATE:
CC PUBLICATION DATE:
CC RELEVANT RESIDUES IN SEQ ID NO:
CC SEQUENCE 21 AA: 2393 MW; 2418 CN;

Query Match 90.1%; Score 136; DB 1; Length 21;
Best Local Similarity 95.2%; Pred. No. 2,71e-08;
Matches 20; Conservative 0; Mismatches 0; Indels 1; Gaps 1

Db 1 SLRPFKALVREKGRHPSHSC 21
|||||
1 SLRPFKALVREKGRHPS-HSC 20

RESULT 3 STANDARD; PRT; 20 AA.
ID US-08-644-456-4
AC xxxxxx
DT
DT
XX Sequence 4, Application US/08644456
DE
CC Sequence 4, Application US/08644456
CC Patent No. 5747650
CC GENERAL INFORMATION:
CC APPLICANT: Kulesz-Martin, Molly F.
CC TITLE OF INVENTION: P53as PROTEIN AND ANTIBODY THEREFOR
CC NUMBER OF SEQUENCES: 5
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Dunn & Associates
CC STREET: P.O. Box 96
CC City: Newfane
CC STATE: New York
CC COUNTRY: U.S.A.
CC ZIP: 14108
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Diskette - 3.50 inch, 1.44 Mb storage
CC COMPUTER: Victor 300 SX/25 (IBM PC Compatible)
CC OPERATING SYSTEM: MS-DOS Version 5.0
CC SOFTWARE: Wordstar Professional Release 4
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/644,456
CC FILING DATE: 10-May-1996
CC CLASSIFICATION: 530
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 08/100,496
CC FILING DATE: 2-Aug-1993
CC CLASSIFICATION: 530
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Dunn, Michael L.
CC REGISTRATION NUMBER: 25,350
CC REFERENCE/DOCKET NUMBER: RPP.135C US
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (716) 433-1661
CC TELEFAX: (716) 433-1665
CC INFORMATION FOR SEQ ID NO: 4:
CC SEQUENCE CHARACTERISTICS:

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CC		LENGTH:	20	
CC		TYPE:	Amino Acids	
CC		STRANDEDNESS:	unknown	
CC		TOPOLOGY:	unknown	
CC		MOLECULE TYPE:	Peptide	
CC		HYPOTHETICAL:	deduced from intron 10 sequences p53 gene	
CC		ANTI-SENSE:		
CC		FRAGMENT TYPE:		
CC		ORIGINAL SOURCE:		
CC		ORGANISM:	Human	
CC		STRAIN:		
CC		INDIVIDUAL ISOLATE:		
CC		DEVELOPMENTAL STAGE:		
CC		HAPLOTYPE:		
CC		TISSUE TYPE:		
CC		CELL TYPE:		
CC		CELL LINE:		
CC		ORGANELLE:		
CC		IMMEDIATE SOURCE:		
CC		LIBRARY:	deduced translation from nucleotides 1n	
CC		LIBRARY:	Genbank nucleic acid database accession #54156,	
CC		LIBRARY:	Locus HSP53G	
CC		CLONE:		
CC		POSITION IN GENOME:		
CC		CHROMOSOME/SEGMENT:	17	
CC		MAP POSITION:	p53 gene, at 18530 to 18589	
CC		UNITS:		
CC		FEATURE:	n/a	
CC		NAME/KEY:		
CC		LOCATION:		
CC		IDENTIFICATION METHOD:		
CC		OTHER INFORMATION:		
CC		PUBLICATION INFORMATION:		
CC		AUTHORS:		
CC		TITLE:		
CC		JOURNAL:		
CC		VOLUME:		
CC		ISSUE:		
CC		PAGES:		
CC		DATE:		
CC		DOCUMENT NUMBER:		
CC		FILING DATE:		
CC		PUBLICATION DATE:		
CC		RELEVANT RESIDUES:	IN SEQ ID NO:	
CC		SEQUENCE	20 AA; 2256 MW; 2309 CN;	
SQ				
		Query Match	56.3%; Score 85; DB 1; Length 20;	
		Best Local Similarity 100.0%; Pred.No. 3.32e-02;		
		Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
DB		1 REKGHRPESHSC 11		
OY		10 REKGHRPESHSC 20		
RESULT	4			
ID	US-07-876-941A-30	STANDARD;	PRT;	18 AA.
XX	xxxxxx			
DT				
XX				
DE	Sequence 30, Application US/07876941A			
CC	Sequence 30, Application US/07876941A			
CC	Patent No. 5885768			
CC	GENERAL INFORMATION:			
CC	APPLICANT: Reyes, Gregory R.			
CC	APPLICANT: Bradley, Daniel W.			
CC	APPLICANT: Tam, Albert W.			
CC	APPLICANT: Mitchell, Carl			
CC	TITLE OF INVENTION: Hepatitis E Virus Peptide Antigen and			
CC	TITLE OF INVENTION: Antibodies			

CC NUMBER OF SEQUENCES: 76
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Dehlinger & Associates
CC STREET: 350 Cambridge Avenue, Suite 250
CC CITY: Palo Alto
CC STATE: CA
CC COUNTRY: USA
CC ZIP: 94306
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Floppy disk
CC COMPUTER: IBM PC compatible
CC OPERATING SYSTEM: PC-DOS/MS-DOS
CC SOFTWARE: Patent Release #1.0, Version #1.25
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/07/876,941A
CC FILING DATE: 01-MAY-1992
CC CLASSIFICATION: 435
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US 822,335
CC FILING DATE: 17-JAN-1992
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US 505,888
CC FILING DATE: 05-APRIL-1990
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US 420,921
CC FILING DATE: 13-OCTOBER-1989
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US 367,486
CC FILING DATE: 16-JUNE-1989
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US 336,672
CC FILING DATE: 11-APRIL-1989
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US 208,997
CC FILING DATE: 17-JUNE-1988
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Sholtz, Charles K.
CC REGISTRATION NUMBER: 38,615
CC REFERENCE/DOCKET NUMBER: 4600-0093.33
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (415) 324-0880
CC TELEFAX: (415) 324-0960
CC INFORMATION FOR SEQ ID NO: 30:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 18 amino acids
CC TYPE: amino acid
CC STRANDEDNESS: single
CC TOPOLOGY: unknown
CC MOLECULE TYPE: peptide
CC HYPOTHETICAL: NO
CC ANTI-SENSE: NO
CC ORIGINAL SOURCE:
CC INDIVIDUAL ISOLATE: FIG. 11, ORF 1, aa 1237-1254
CC SEQUENCE 18 AA; 2035 MW; 1618 CN;
SQ
Query Match 33.1%; Score 50; DB 2; Length 18;
Best Local Similarity 46.7%; Pired. NO. 1.85e+02;
Matches 7; Conservative 2; Mismatches 6; Indels 0; Gaps 0;
DB 3 ISAFHOLAEELGHRP 17
QY 2 LRFKALVREKGRHP 16

RESULT 5
ID US-08-644-456-1 STANDARD; PRT: 17 AA.
XX xxxxxx
XX
XX
XX
XX Sequence 1, Application US/08644456

CC Sequence 1, Application US/08644456
CC Patent No. 5747650
CC GENERAL INFORMATION:
CC APPLICANT: Kulesz-Martin, Molly F.
CC TITLE OF INVENTION: p53as PROTEIN AND ANTIBODY THEREFOR
CC NUMBER OF SEQUENCES: 5
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Dunn & Associates
CC STREET: P.O. Box 96
CC CITY: Newtane
CC STATE: New York
CC COUNTRY: U.S.A.
CC ZIP: 14108
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: Diskette - 3.50 inch, 1.44 Mb storage
CC COMPUTER: VICTOR 300 SX/25 (IBM PC Compatible)
CC OPERATING SYSTEM: MS-DOS Version 5.0
CC SOFTWARE: Wordstar Professional Release 4
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/644,456
CC FILING DATE: 10-May-1996
CC CLASSIFICATION: 530
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 08/100,496
CC FILING DATE: 2-Aug-1993
CC CLASSIFICATION: 530
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Dunn, Michael L.
CC REGISTRATION NUMBER: 25,330
CC REFERENCE/DOCKET NUMBER: RPP:135C US
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (716) 433-1661
CC TELEFAX: (716) 433-1665
CC INFORMATION FOR SEQ ID NO: 1:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 17
CC TYPE: amino acids
CC STRANDEDNESS: n/a
CC TOPOLOGY: n/a
CC MOLECULE TYPE: peptide
CC HYPOTHETICAL: no
CC ANTI-SENSE: no
CC FRAGMENT TYPE: n/a
CC ORIGINAL SOURCE:
CC ORGANISM: mouse
CC STRAIN: n/a
CC INDIVIDUAL ISOLATE: n/a
CC DEVELOPMENTAL STAGE: n/a
CC HAPLOTYPE: n/a
CC TISSUE TYPE: n/a
CC CELL LINE: n/a
CC ORGANELLE: n/a
CC IMMEDIATE SOURCE: sequenced from cDNA clone from mouse
CC IMMEDIATE SOURCE: epidermal cell RNA, Genbank Accession #M13874
CC LIBRARY: plasmid p6.3
CC POSITION IN GENOME:
CC CHROMOSOME/SEGMENT: 11
CC MAP POSITION: p53 gene
CC UNITS:
CC FEATURE: n/a
CC NAME/KEY:
CC LOCATION:
CC IDENTIFICATION METHOD:
CC OTHER INFORMATION:
CC PUBLICATION INFORMATION:
CC AUTHORS: Kulesz-Martin et al.
CC TITLE: Endogenous p53 Protein Generated from Wild
CC JOURNAL: Mol. Cell. Biol.
CC VOLUME: 14
CC ISSUE: 3

CC PAGES: 1698-1708
CC DATE: March, 1994
CC AUTHORS: Han, K.A. and Kulesz-Martin, M.F.
CC TITLE: Alternatively Spliced p53 RNA in Transformed
CC TITLE: and No. 5747650mal Cells of Different Tissue Types
CC JOURNAL: Nucleic Acids Res.
CC VOLUME: 20
CC ISSUE: 8
CC PAGES: 1979-1981
CC DATE: 1992
CC AUTHORS: Aral, N. et al.
CC TITLE: Immunologically Distinct p53 Molecules Generated
CC TITLE: by Alternative Splicing
CC JOURNAL: Mol. and Cell. Biol.
CC VOLUME: 6
CC ISSUE: 6
CC PAGES: 3232-3239
CC DATE: 1986
CC DOCUMENT NUMBER:
CC FILING DATE:
CC PUBLICATION DATE:
CC RELEVANT RESIDUES IN SEQ ID NO:
SQ SEQUENCE 17 AA; 1944 MW; 1303 CN;

Query Match 31.8%; Score 48; DB 1; Length 17;
Best Local Similarity 55.6%; Pred. No. 2.89e+02;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Db 4 RAFOALIKE 12
| | | | |
QY 3 RPFKALVRE 11

RESULT 6
ID US-08-313-681A-11 STANDARD; PRT; 16 AA.
XX xxxxxx

Sequence 11, Application US/08313681A
Patent No. 5618675
GENERAL INFORMATION:
APPLICANT: Larrick, James W.
APPLICANT: Wright, Susan C.
APPLICANT: Hirata, Mishima
TITLE OF INVENTION: Human Cationic Proteins Having
TITLE OF INVENTION: Lipopolysaccharide Binding and Anti-Coagulant Activity
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend Kourile and Crew
STREET: One Market Plaza, Stewart Tower, Suite 2000
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/313,681A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Heslin, James M.
REGISTRATION NUMBER: 29,541
REFERENCE/DOCKET NUMBER: 15325-9-1
TELEPHONE: 415-326-2400

CC TELEFAX: 415-326-2422
CC INFORMATION FOR SEQ ID NO: 11:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 16 amino acids
CC TYPE: amino acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
CC MOLECULE TYPE: peptide
SQ SEQUENCE 16 AA; 2071 MW; 1188 CN;

Query Match 30.5%; Score 46; DB 1; Length 16;
Best Local Similarity 45.5%; Pred. No. 4.48e+02;
Matches 5; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Db 6 LRFRNKIKER 16
| | | | |
QY 2 LRPFKALVREK 12

RESULT 7
ID US-08-313-681A-7 STANDARD; PRT; 29 AA.
XX xxxxxx

Sequence 7, Application US/08313681A
Patent No. 5618675
GENERAL INFORMATION:
APPLICANT: Larrick, James W.
APPLICANT: Wright, Susan C.
APPLICANT: Hirata, Mishima
TITLE OF INVENTION: Human Cationic Proteins Having
TITLE OF INVENTION: Lipopolysaccharide Binding and Anti-Coagulant Activity
NUMBER OF SEQUENCES: 30
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend Kourile and Crew
STREET: One Market Plaza, Stewart Tower, Suite 2000
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94105
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/313,681A
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Heslin, James M.
REGISTRATION NUMBER: 29,541
REFERENCE/DOCKET NUMBER: 15325-9-1
TELEPHONE: 415-326-2400
TELEFAX: 415-326-2422
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 29 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Region
LOCATION: 23
OTHER INFORMATION: /note= "Xaa is Asp or Lys"
FEATURE:
NAME/KEY: Region

CC LOCATION: 26
CC OTHER INFORMATION: /note- "Xaa is a Gln or Ile"
CC FEATURE:
CC NAME/KEY: Region
CC LOCATION: 27
CC OTHER INFORMATION: /note- "Xaa is a Gly or Gln"
CC SEQUENCE 29 AA: 3536 MW; 5084 CN;
SQ

Query Match 30.5%; Score 46; DB 1; Length 29;
Best Local Similarity 45.5%; Pred. No. 4,486+02;
Matches 5; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Db 6 LRRFRNKIKR 16
|||: |||
QY 2 LRRFKALVRK 12

RESULT 8
ID US-08-188-583-32 STANDARD; PRT; 12 AA.
XX xxxxxx

Sequence 32, Application US/08188583
CC Patent No. 5851813
CC GENERAL INFORMATION:
CC APPLICANT: Desrosiers, Ronald C.
CC TITLE OF INVENTION: PRIMATE LENTIVIRUS VACCINES
CC NUMBER OF SEQUENCES: 57
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: Fish & Richardson
CC STREET: 225 Franklin Street
CC CITY: Boston
CC STATE: Massachusetts
CC COUNTRY: U.S.A.
CC ZIP: 02110-2804
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
CC OPERATING SYSTEM: IBM PS/2 Model 502 or 555X
CC SOFTWARE: WordPerfect (Version 5.0)
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/188,583
CC FILING DATE:
CC CLASSIFICATION: 435
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 07/727,494
CC FILING DATE: July 9, 1991
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 07/551,945
CC FILING DATE: July 12, 1990
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Freeman, John W.
CC REGISTRATION NUMBER: Reg. No. 5851813 29,066
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (617) 542-5070
CC TELEFAX: (617) 542-8906
CC TELEX: 200154
CC INFORMATION FOR SEQ ID NO: 32:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 12
CC TYPE: amino acid
CC STRANDEDNESS:
CC TOPOLOGY: linear
SQ SEQUENCE 12 AA: 1318 MW; 741 CN;

Query Match 29.8%; Score 45; DB 2; Length 12;
Best Local Similarity 75.0%; Pred. No. 5,566+02;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Db 1 KGRGSH 8
|||: |||
QY 12 KGRPSHS 19

RESULT 9
ID US-08-232-453A-41 STANDARD; PRT; 14 AA.
XX xxxxxx

Sequence 41, Application US/08232453A
CC Patent No. 5589568
CC GENERAL INFORMATION:
CC APPLICANT: HIGASHIJIMA, TSUTOMU
CC APPLICANT: ROSS, ELIOTT M.
CC TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
CC TITLE OF INVENTION: MODULATING G PROTEIN ACTION
CC NUMBER OF SEQUENCES: 71
CC CORRESPONDENCE ADDRESS:
CC ADDRESSEE: ARNOLD, WHITE & DURKEE
CC STREET: P.O. BOX 4433
CC CITY: HOUSTON
CC STATE: TX
CC COUNTRY: USA
CC ZIP: 77210
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: FLOPPY DISK
CC OPERATING SYSTEM: IBM PC COMPATIBLE
CC SOFTWARE: WORDPERFECT 5.1
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/232,453A
CC FILING DATE: APRIL 22, 1994
CC CLASSIFICATION: 514
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: US 07/748,319
CC FILING DATE: AUGUST 21, 1991
CC CLASSIFICATION: 514
CC ATTORNEY/AGENT INFORMATION:
CC NAME: PARKER, DAVID L.
CC REGISTRATION NUMBER: 32,165
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (512) 418-3000
CC TELEFAX: (512) 474-7577
CC INFORMATION FOR SEQ ID NO: 41:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 14 amino acids
CC TYPE: amino acid
CC STRANDEDNESS: single
CC TOPOLOGY: linear
SQ SEQUENCE 14 AA: 1564 MW; 590 CN;

Query Match 29.8%; Score 45; DB 1; Length 14;
Best Local Similarity 50.0%; Pred. No. 5,566+02;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Db 2 NRALRALAR 11
|||: |||
QY 1 SLRPFKALVR 10

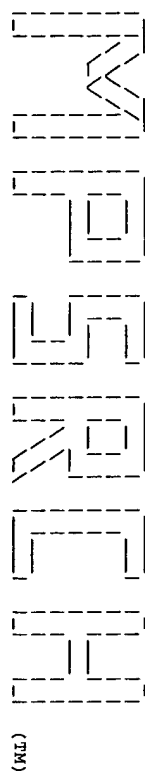
RESULT 10
ID US-07-603-782A-1 STANDARD; PRT; 27 AA.
XX xxxxxx

CC	APPLICATION NUMBER:	US/08/224,6532A
CC	FILING DATE:	
CC	CLASSIFICATION:	530
CC	ATTORNEY/AGENT INFORMATION:	
CC	NAME:	Goltick, Mary E.
CC	REGISTRATION NUMBER:	34,829
CC	REFERENCE/DOCKET NUMBER:	22727-00111
CC	TELECOMMUNICATION INFORMATION:	
CC	TELEPHONE:	216 622 8458
CC	TELEFAX:	216 241 0816
CC	INFORMATION FOR SEQ ID NO:	4:
CC	SEQUENCE CHARACTERISTICS:	
CC	LENGTH:	15 amino acids
CC	TYPE:	amino acid
CC	STRANDEDNESS:	
CC	TOPOLOGY:	linear
CC	MOLECULE TYPE:	peptide
CC	HYPOTHETICAL:	NO
CC	SEQUENCE	15 AA; 1756 MW; 1288 CN;
DB	6 OKHGRGTH 13	
	: :1	
QY	11 EKGRPSH 18	
RESULT	13	
ID	US-07-882-923-7	STANDARD; PRT; 20 AA.
XX	xxxxxx	
XX		
DT		
XX		
DE	Sequence 7, Application US/07882923	
XX		
CC	Sequence 7, Application US/07882923	
CC	Patent No. 5328899	
CC	GENERAL INFORMATION:	
CC	APPLICANT:	Boublik, Jaroslav H.
CC	APPLICANT:	Rivier, Jean E.F.
CC	APPLICANT:	Brown, Marvin R.
CC	APPLICANT:	Scott, Neal A.
CC	TITLE OF INVENTION:	NPY PEPTIDE ANALOGS
CC	NUMBER OF SEQUENCES:	14
CC	CORRESPONDENCE ADDRESSES:	
CC	ADDRESSEE:	Fitch, Even, Tabin & Flannery
CC	STREET:	4250 Executive Square, Suite 510
CC	CITY:	La Jolla
CC	STATE:	CA
CC	COUNTRY:	USA
CC	ZIP:	92037
CC	COMPUTER READABLE FORM:	
CC	MEDIUM TYPE:	Floppy disk
CC	COMPUTER:	IBM PC compatible
CC	OPERATING SYSTEM:	PC-DOS/MS-DOS
CC	SOFTWARE:	Patentin Release #1.0, Version #1.25
CC	CURRENT APPLICATION DATA:	
CC	APPLICATION NUMBER:	US/07/882,923
CC	FILING DATE:	19920512
CC	CLASSIFICATION:	514
CC	PRIOR APPLICATION DATA:	
CC	APPLICATION NUMBER:	US 07/503,198
CC	FILING DATE:	30-MAR-1990
CC	PRIOR APPLICATION DATA:	
CC	APPLICATION NUMBER:	US 07/219,596
CC	FILING DATE:	15-JUL-1988
CC	ATTORNEY/AGENT INFORMATION:	
CC	NAME:	Schumann, James J
CC	REGISTRATION NUMBER:	20,856
CC	REFERENCE/DOCKET NUMBER:	52864

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CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: 619-552-1311
CC TELEFAX: 619-552-0095
CC INFORMATION FOR SEQ. ID NO: 7:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 20 amino acids
CC TYPE: AMINO ACID
CC TOPOLOGY: unknown
CC MOLECULE TYPE: peptide
CC SEQUENCE 20 AA: 2589 MW; 1767 CN;
SQ
    Query Match          29.1%; Score 44; DB 1; Length 20;
    Best Local Similarity 30.8%; Pred. No. 6,88e+02;
    Matches 4; Conservative 6; Mismatches 2; Indels 1; Gaps 1;
Db
    7 ALROYNLITROR 19
    :|:|:|:|:|:|:
Oy 1 SLRPFKALY-REK 12
RESULT 14
ID US-08-329-151-28 STANDARD; PRT; 15 AA.
XX xxxxxx
XX
XX
XX
DE Sequence 28, Application US/08329151
CC
CC Sequence 28, Application US/08329151
CC Patent No. 5604203
CC GENERAL INFORMATION:
CC APPLICANT: Balasubramaniam, A.
CC TITLE OF INVENTION: ANALOGS OF PEPTIDE YY AND USES
CC TITLE OF INVENTION: THEREOF
CC NUMBER OF SEQUENCES: 30
CC CORRESPONDENCE ADDRESSES:
CC ADDRESSEE: Fish & Richardson
CC STREET: 225 Franklin Street
CC CITY: Boston
CC STATE: Massachusetts
CC COUNTRY: U.S.A.
CC ZIP: 02110-2804
CC
CC COMPUTER READABLE FORM:
CC MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
CC COMPUTER: IBM PS/2 Model 50Z or 55SX
CC OPERATING SYSTEM: MS-DOS (Version 5.1)
CC SOFTWARE: Wordperfect (Version 5.1)
CC CURRENT APPLICATION DATA:
CC APPLICATION NUMBER: US/08/329,151
CC FILING DATE:
CC
CC CLASSIFICATION: 514
CC PRIOR APPLICATION DATA:
CC APPLICATION NUMBER: 08/038,534
CC FILING DATE: 3/29/93
CC APPLICATION NUMBER: 08/109,326
CC FILING DATE: 08/19/93
CC ATTORNEY/AGENT INFORMATION:
CC NAME: Paul T. Clark
CC REGISTRATION NUMBER: 30,162
CC REFERENCE/DOCKET NUMBER: 00537/105001
CC TELECOMMUNICATION INFORMATION:
CC TELEPHONE: (617) 542-5070
CC TELEFAX: (617) 542-8906
CC TELEX: 200154
CC INFORMATION FOR SEQ. ID NO: 28:
CC SEQUENCE CHARACTERISTICS:
CC LENGTH: 15
CC TYPE: amino acid
CC STRANDEDNESS: N/A
CC TOPOLOGY: linear
CC FEATURE:
CC OTHER INFORMATION: xaa in position 5 is an abbreviation of

```

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MPsrch_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Tue Jun 22 14:56:29 1999; Maspar time 4.30 Seconds
186.302 Million cell updates/sec

Tabular output not generated.

Title: >US-08-644-289-1
Description: (1-20) from US08644289.pep
Perfect Score: 151
Sequence: 1 SLRPFKALVREKGRPSHSC 20

Scoring table: PAM 150
Gap 15

Searched: 122810 seqs, 40068593 residues

Post-processing: Minimum Match 08
Listing first 1000 summaries
Maximum DB seq length 30

Database: p1r60
1:p1r1 2:p1r2 3:p1r3 4:p1r4

Statistics: Mean 28.488; Variance 40.935; scale 0.696

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query					
No.	Score	Match	Length	ID	Description	Pred. No.
-----	-----	-----	-----	-----	-----	-----

No matches found.

Search completed: Tue Jun 22 14:57:18 1999
Job time : 49 secs.



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Msearch_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Tue Jun 22 14:57:36 1999; MasPar time 2.77 Seconds
203.787 Million cell updates/sec

Tabular output not generated.

Title: >US-08-644-289-1
Description: (1-20) from US08644289.pep
Perfect Score: 151
Sequence: 1 SLRPFKALVREKGRPSHSC 20

Scoring table:
PAM 150
Gap 15

Searched: 77977 seqs, 28268293 residues

Post-processing: Minimum Match 0%
Listing first 1000 summaries
Maximum DB seq length 30

Database: swiss-prot37
1:swissprot

Statistics: Mean 29.227; Variance 35.378; scale 0.826







Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query	No.	Score	Match	Length	ID	Description	Pred.	No.
--------	-------	-----	-------	-------	--------	----	-------------	-------	-----

No matches found.

Search completed: Tue Jun 22 14:58:30 1999
Job time : 54 secs.

(TM)

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```

MPsrch_pp  protein - protein database search, using Smith-Waterman algorithm
Run on:      Tue Jun 22 14:58:48 1999;  MasPar time 5.55 Seconds
Tabular output not generated.  196.656 Million cell updates/sec

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Title:	>US-08-644-289-1
Description:	(1-20) from US08644289.pep
Perfect Score:	151
Sequence:	1 SLRPFKALVREKGHRPSHSC 20

Scoring table: PAM 150
Gap 15

Searched: 179066 seqs, 54579741 residues

```
Post-processing: Minimum Match 0%
                  Listing first 1000 summaries
                  Maximum DB seq length 30
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Database: **sptrembl9**
1:sp_archaea 2:sp_bacteria 3:sp_fungi 4:sp_human
5:sp_invertebrate 6:sp_mammal 7:sp_mhc 8:sp_organelle
9:sp_phage 10:sp_plant 11:sp_podent 12:sp_unclassified
13:sp_vertebrate 14:sp_virus

Statistics: Mean 28.060; Variance 36.106; scale 0.777

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

SUMMARIES

Result	Query	ID	Description	Pred. No.
No.	Score	Match	Length	DB

No matches found.

Search completed: Tue Jun 22 15:00:25 1999
Job time : 97 secs.